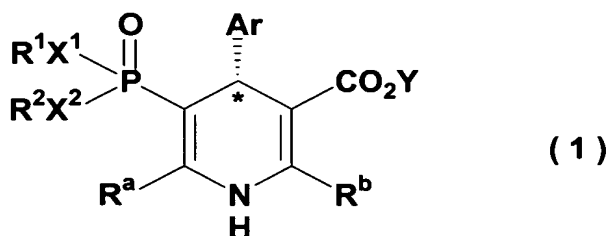


Amendments to the Claims:

The following listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) A T-type calcium channel blocker that is optically active 1,4-dihydropyridine compound, a pharmaceutically acceptable salt thereof or a solvate thereof, of formula (1)



[wherein

R^1 and R^2 are independently of each other C_{1-6} alkyl group {the C_{1-6} alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom), C_{2-6} alkenyl group or C_{2-6} alkynyl group (the C_{2-6} alkenyl group and C_{2-6} alkynyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom))}, or $-L^1-NR^3R^4$ { R^3 and R^4 are independently of each other C_{1-6} alkyl group (the C_{1-6} alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom)) or phenyl group (wherein the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom), L^1 is C_{2-6} alkylene group (the C_{2-6} alkylene group may be substituted with C_{1-3} alkyl group or phenyl group (the phenyl group may be arbitrarily substituted with halogen atom, C_{1-3} alkyl group or C_{1-3} alkoxy group))}, or

R^1 and R^2 together form $-CR^5R^6-CR^7R^8-$, $-CR^5R^6-CR^7R^8-CR^9R^{10}-$ or $-CR^5R^6-CR^7R^8-CR^9R^{10}-CR^{11}R^{12}-$ (R^5 to R^{12} are independently of each other hydrogen atom or C_{1-6} alkyl

group, or any two of them together with the carbon atom bonding them may form 5-, 6- or 7-membered ring);

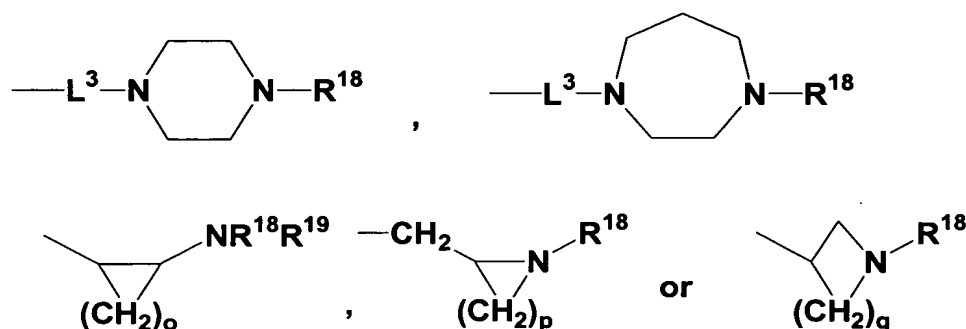
X^1 and X^2 are independently of each other O or NR^{13} (R^{13} is hydrogen atom or C_{1-6} alkyl group);

Ar is phenyl group, pyridyl group, furyl group or 2,1,3-benzoxadiazol-4-yl group {the phenyl group, pyridyl group, furyl group and 2,1,3-benzoxadiazol-4-yl group may arbitrarily substituted with one or two substituents selected from NO_2 , CF_3 , Br, Cl, F, R (R is C_{1-20} alkyl group), OH, OR^{14} (R^{14} is C_{1-6} alkyl group), $OCHF_2$, $COOR^{14}$, NH_2 , NHR^{14} , $NR^{14}R^{15}$ (R^{15} is C_{1-6} alkyl group), $CONH_2$, $CONHR^{14}$, $CONR^{14}R^{15}$, $COSR^{14}$, SR^{14} , $S(O)R^{14}$, $S(O)_2R^{14}$, SO_3H , SO_3R^{14} , SO_2NH_2 , SO_2NHR^{14} , $SO_2NR^{14}R^{15}$, CN and phenyloxy group};

R^a and R^b are independently of each other C_{1-6} alkyl group, $-L^2-NR^{16}R^{17}$ { R^{16} and R^{17} are independently of each other hydrogen atom, C_{1-6} alkyl group (the C_{1-6} alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom)) or phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom)}, L^2 is C_{2-6} alkylene group (the C_{2-6} alkylene group may be arbitrarily substituted with C_{1-3} alkyl group or phenyl group (the phenyl group may be arbitrarily substituted with halogen atom, C_{1-3} alkyl group or C_{1-3} alkoxy group))), $CH_2O-L^2-NR^{16}R^{17}$, Ar^1 (Ar^1 is phenyl group (the phenyl group may be arbitrarily substituted with halogen atom, C_{1-3} alkyl group or C_{1-3} alkoxy group)), $CH=CHAr^1$, $CH_2CH(OH)Ar^1$, CHO, CN, CH_2OH , $CHOR^{16}$, $-L^2-N(CH_2CH_2)_2NR^{16}$ or $NR^{16}R^{17}$;

Y is C_{1-20} alkyl group {the C_{1-20} alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom), C_{2-6} alkenyl group or C_{2-6} alkynyl group (the C_{2-6} alkenyl group and C_{2-6} alkynyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom))}, $-L^3-NR^{18}R^{19}$ { R^{18} and R^{19} are independently of each other

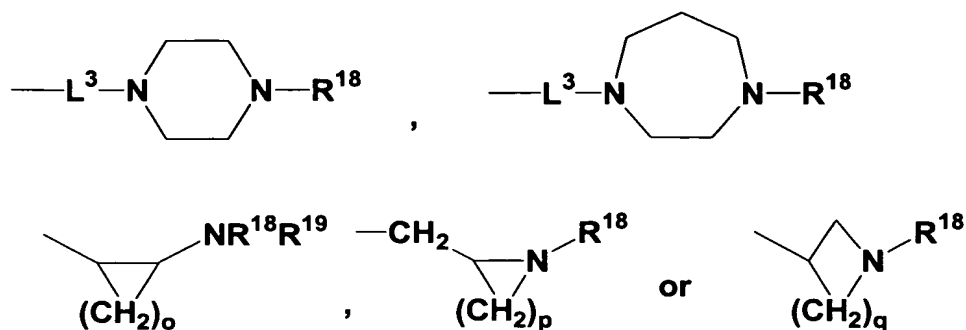
C₁₋₆ alkyl group (the C₁₋₆ alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C₁₋₆ alkoxy group or halogen atom)) or phenyl group (the phenyl group may be substituted with C₁₋₆ alkoxy group or halogen atom), L³ is C₂₋₆ alkylene group (the C₂₋₆ alkylene group may be arbitrarily substituted with C₁₋₃ alkyl group or phenyl group (the phenyl group may be arbitrarily substituted with halogen atom, C₁₋₃ alkyl group or C₁₋₃ alkoxy group))),



(wherein o and p are independently of each other 3 or 4, q is 1, 2 or 3), and

* is absolute configuration of R.]

2. (Original) The T-type calcium channel blocker that is optically active 1,4-dihydropyridine compound, a pharmaceutically acceptable salt thereof or a solvate thereof, according to claim 1, wherein Y is -L³-NR¹⁸R¹⁹ {R¹⁸ and R¹⁹ are independently of each other C₁₋₆ alkyl group (the C₁₋₆ alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C₁₋₆ alkoxy group or halogen atom)) or phenyl group (the phenyl group may be substituted with C₁₋₆ alkoxy group or halogen atom), L³ is C₂₋₆ alkylene group (the C₂₋₆ alkylene group may be arbitrarily substituted with C₁₋₃ alkyl group or phenyl group (the phenyl group may be arbitrarily substituted with halogen atom, C₁₋₃ alkyl group or C₁₋₃ alkoxy group))),



(wherein o and p are independently of each other 3 or 4, q is 1, 2 or 3), and

R^a is C_{1-6} alkyl group.

3. (Original) The T-type calcium channel blocker that is optically active 1,4-dihydropyridine compound, a pharmaceutically acceptable salt thereof or a solvate thereof, according to claim 2, wherein R^b is C_{1-6} alkyl group, CN or NH_2 .

4. (Original) The T-type calcium channel blocker that is optically active 1,4-dihydropyridine compound, a pharmaceutically acceptable salt thereof or a solvate thereof, according to claim 1, wherein Y is C_{1-20} alkyl group {the C_{1-20} alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom), C_{2-6} alkenyl group or C_{2-6} alkynyl group (the C_{2-6} alkenyl group and C_{2-6} alkynyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom))},

R^b is $-L^2-NR^{16}R^{17}$ { R^{16} and R^{17} are independently of each other hydrogen atom, C_{1-6} alkyl group (the C_{1-6} alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom)) or phenyl group (the phenyl group may be substituted with C_{1-6} alkoxy group or halogen atom)}, L^2 is C_{2-6} alkylene group (the C_{2-6} alkylene group may be arbitrarily substituted with C_{1-3} alkyl group or phenyl group (the

phenyl group may be arbitrarily substituted with halogen atom, C₁₋₃ alkyl group or C₁₋₃ alkoxy group))), CH₂O-L²-NR¹⁶R¹⁷ or -L²-N(CH₂CH₂)₂NR¹⁶, and R^a is C₁₋₆ alkyl group.

5. (Currently Amended) The T-type calcium channel blocker that is optically active 1,4-dihydropyridine compound, a pharmaceutically acceptable salt thereof or a solvate thereof, according to claim 2, ~~3 or 4~~, wherein R¹ and R² are independently of each other C₁₋₆ alkyl group {the C₁₋₆ alkyl group may be substituted with phenyl group (the phenyl group may be substituted with C₁₋₆ alkoxy group or halogen atom), C₂₋₆ alkenyl group or C₂₋₆ alkynyl group (the C₂₋₆ alkenyl group and C₂₋₆ alkynyl group may be substituted with phenyl group (the phenyl group may be substituted with C₁₋₆ alkoxy group or halogen atom))}, or R¹ and R² together form -CR⁵R⁶-CR⁷R⁸-, -CR⁵R⁶-CR⁷R⁸-CR⁹R¹⁰- or -CR⁵R⁶-CR⁷R⁸-CR⁹R¹⁰-CR¹¹R¹²- (R⁵ to R¹² are independently of each other hydrogen atom or C₁₋₆ alkyl group, or any two of them together with the carbon atom bonding them may form 5-, 6- or 7-membered ring); X¹ and X² are both O.

6. (Original) The T-type calcium channel blocker that is optically active 1,4-dihydropyridine compound, a pharmaceutically acceptable salt thereof or a solvate thereof, according to claim 5, wherein Ar is phenyl, 3-nitrophenyl, 2-nitrophenyl, 3-chlorophenyl, 2-chlorophenyl, 3-methoxyphenyl, 2-methoxyphenyl, 2-trifluoromethylphenyl, 2-trifluoromethylphenyl or 2,3-dichlorophenyl.

7. (Original) The T-type calcium channel blocker that is optically active 1,4-dihydropyridine compound, a pharmaceutically acceptable salt thereof or a solvate thereof, according to claim 6, wherein R¹ and R² together form -CH₂-C(CH₃)₂-CH₂-, X¹ and X² are

both O, Ar is 3-nitrophenyl, R^a and R^b are both methyl, and Y is 2-[benzyl(phenyl)amino]ethyl.

8. (Currently Amended) A pharmaceutical containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

9. (Currently Amended) A therapeutic or preventive agent against a disease for which T-type calcium channel blocking action is effective, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

10. (Currently Amended) A therapeutic or preventive agent against hypercardia, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

11. (Currently Amended) A therapeutic or preventive agent against heart failure, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

12. (Currently Amended) A therapeutic or preventive agent against cardiomyopathy, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

13. (Currently Amended) A therapeutic or preventive agent against atrial fibrillation, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

14. (Currently Amended) A therapeutic or preventive agent against tachycardia-arrhythmia, containing the T-type calcium channel blocker according to ~~any one of claims 1~~

~~to 7~~claim 1.

15. (Currently Amended) A therapeutic or preventive agent against arterial sclerosis, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

16. (Currently Amended) A therapeutic or preventive agent against nephritis, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

17. (Currently Amended) A therapeutic or preventive agent against nephropathy, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

18. (Currently Amended) A therapeutic or preventive agent against renal disorder, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

19. (Currently Amended) A therapeutic or preventive agent against renal insufficiency, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

20. (Currently Amended) A therapeutic or preventive agent against edema, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

21. (Currently Amended) A therapeutic or preventive agent against inflammation, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

22. (Currently Amended) A therapeutic or preventive agent against hyperaldosteronism, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

23. (Currently Amended) A therapeutic or preventive agent against neurogenic pain, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.

24. (Currently Amended) A therapeutic or preventive agent against epilepsy, containing the T-type calcium channel blocker according to ~~any one of claims 1 to 7~~claim 1.